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				Date of Report PRISCA	8/6/2023 5.1.0.17
Patient Data					
Name	MRS. PARA	MA HALDER	Patient ID		012306060219
Birthday		19/10/1998	Sample ID		11571010
Age at term		25	Sample Date		6/6/2023
Gestational age		13+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57.9 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
<b>Biochemical Data</b>			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	13+4
PAPP-A	4.32 mIU/ml	0.53	Method		CRL (<>Robinson)
fb-hCG	34.1 ng/ml	1.23	Scan date		6/6/2023
Risks at sampling date			Nuchal translucency (NT) 1.8		
Age Risk 1:1010		1:1010	Nuchal translucency MoM 0.		0.97
Biochemical T21 risk 1:84		1:844	Nasal bone		present
Combined trisomy 21 risk	ζ.	1:3966			
Trisomy 13/18		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
Risk 1:10 1:250 Cut off 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test it is expected that among 3966 women with the same data, there is one woman with a trisomy 21 pregnancy and 3965 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		